

Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

1. **(Currently Amended)** A method for forming a two-dimensional ordered array of proteins, comprising:
 - contacting a population of proteins with a gas-aqueous interface;
 - laterally compressing said population to an appropriate pressure, such that a two-dimensional ordered array of said proteins is formed at said interface, wherein said proteins are not solubilized using detergent.
 2. **(Cancelled).**
 3. **(Previously Presented)** The method of claim 64, wherein said amphiphilic molecule comprises a protein.
 4. **(Previously Presented)** The method of claim 1, wherein said protein is a membrane protein, a cellular receptor, an orphan receptor, receptor tyrosine kinase, an EPH receptor, an ion channel, a cytokine receptor, an multisubunit immune recognition receptor, a chemokine receptor, a growth factor receptor, or a G-protein coupled receptor.
 5. **(Previously Presented)** The method of claim 1, wherein said protein is contacted with said interface in the presence of lipids.
 6. **(Previously Presented)** The method of claim 1, further comprising applying said proteins to said interface in proteoliposomes, liposomes, or a cellular membrane.
 7. **(Cancelled).**
 8. **(Currently Amended)** The method of claim 1, wherein said interface is an air-aqueous interface.
- Claims 9-62 **(Cancelled).**

63. **(Currently Amended)** A method for forming a two- or three-dimensional ordered array of water insoluble membrane proteins, comprising:

contacting a population of water insoluble membrane proteins with a gas-aqueous interface, wherein said population of membrane proteins are applied to said interface in a proteoliposome;

laterally compressing said population to an appropriate pressure, such that a two- or three-dimensional ordered array of said water insoluble membrane proteins is formed at said gas-aqueous interface.

64. **(Currently Amended)** A method for forming a three-dimensional ordered array of ~~amphiphilic molecules~~ water insoluble membrane proteins, comprising:

contacting a population of ~~amphiphilic molecules~~ water insoluble membrane proteins with a gas-aqueous interface;

laterally compressing said population to an appropriate pressure, such that a three-dimensional ordered array of said ~~amphiphilic molecules~~ water insoluble membrane proteins is formed at said interface, wherein said appropriate pressure is above a critical density point for the formation of a two-dimensional ordered array of said ~~amphiphilic molecules~~ water insoluble membrane proteins.

Claims 65-66. **(Cancelled)**.

67. **(Previously Presented)** The method of claim 1, wherein said two-dimensional ordered array is a two-dimensional crystalline array.

68. **(Previously Presented)** The method of claim 64, wherein said three-dimensional ordered array is a three-dimensional crystalline array.

69. **(Previously Presented)** The method of claim 3, wherein said protein is a membrane protein, a cellular receptor, an orphan receptor, receptor tyrosine kinase, an EPH receptor, an ion channel, a cytokine receptor, an multisubunit immune recognition receptor, a chemokine receptor, a growth factor receptor, or a G-protein coupled receptor.

70. **(Previously Presented)** The method of claim 3, wherein said protein is contacted with said interface in the presence of lipids.

71. **(Previously Presented)** The method of claim 3, further comprising applying said proteins to said interface in proteoliposomes, liposomes, or a cellular membrane.

Claims 72-73 **(Cancelled)**.

74. **(New)** A method for forming a two- or three- dimensional ordered array of proteins suitable for use in crystallography to determine said protein's structure, comprising:
contacting a population of proteins with a gas-aqueous interface;
laterally compressing said population to an appropriate pressure, such that a two-dimensional ordered array of said proteins is formed at said interface, wherein the structure of said protein using said two- or three- dimensional ordered array can be determined to a resolution of 5 Å or higher.

75. **(New)** A method for forming a two-dimensional ordered array of proteins, comprising:
contacting a population of proteins with a gas-aqueous interface;
laterally compressing said population to an appropriate pressure, such that a two-dimensional ordered array of said proteins is formed at said interface, wherein said two-dimensional ordered array is formed in the absence of a ligand of said protein.

76. **(New)** A method for forming a two- or three-dimensional ordered array of water insoluble membrane proteins, comprising:
contacting a population of water insoluble membrane proteins with a gas-aqueous interface;
laterally compressing said population to an appropriate pressure, such that a two- or three-dimensional ordered array of said water insoluble membrane proteins is formed at said gas-aqueous interface.